

Medication Coverage Exception

| Member and Medication Information (required) | | |
|---|---------------------|-------------|
| Member ID: | Member Name: | |
| DOB: | Weight: | |
| Medication Name/ Strength: | Dose: | |
| <input type="checkbox"/> Do Not Substitute. Authorizations will be processed for the preferred Generic/Brand equivalent unless otherwise specified. | Directions for use: | |
| Provider Information (required) | | |
| Name: | NPI: | Specialty: |
| Contact Person: | Office Phone: | Office Fax: |
| FAX FORM AND RELEVANT DOCUMENTATION INCLUDING: LABORATORY RESULTS, CHART NOTES and/or UPDATED PROVIDER LETTER TO 855-828-4992 | | |

Please select which type of prior authorization you are requesting (check all that apply):

- | | | | |
|--|--|--|-------------------------------------|
| <input type="checkbox"/> Non-preferred | <input type="checkbox"/> Brand Name | <input type="checkbox"/> Combination Product | <input type="checkbox"/> Dosing Kit |
| <input type="checkbox"/> Off-Label Use | <input type="checkbox"/> Limit Exception | <input type="checkbox"/> Step Therapy | <input type="checkbox"/> Other |

Non-Preferred Criteria for Approval: *(at least one of the following conditions must be met)*

- ☐ Trial and failure at an appropriate dose and duration of at least one preferred agent in the drug class.
 Medication and dose: _____ Chart Note Page #: _____
 Details of failure: _____
- ☐ Appropriate clinical rationale for prescribing the non-preferred product: *(adverse reaction, allergy, or inadequate response)*
 _____ Chart Note Page #: _____
- ☐ Continuation of Therapy: Member has been treated with the requested non-preferred drug at a consistent dosage for at least 60 days in most recent 90 days and the prescriber indicates the prescribed medication will best treat the member's condition. Details of therapy (including dates): _____
 _____ Chart Note Page #: _____

Brand Name Medication Criterion for Approval:

- ☐ Appropriate clinical rationale for dispensing the brand name medication:
 _____ Chart Note Page #: _____

Combination Product Criteria for Approval: *Utah Medicaid requires the use of multiple single-entity products instead of one combination product, unless the combination is listed as preferred on the Utah Medicaid Preferred Drug List.*

- ☐ Trial and failure of individual agents in the combination product OR trial and failure of a preferred agent in each of the combination product's therapeutic drug classes.
 Medications used: _____ Chart Note Page #: _____
 Details of failure: _____
- ☐ Appropriate clinical rationale for prescribing the combination product:
 _____ Chart Note Page #: _____

Dosing Kit Criteria for Approval: *Utah Medicaid does not reimburse for dosing kits (e.g. therapy initiation dose titration kits), unless a product is only available in a kit.*

- ☐ Appropriate clinical rationale for prescribing the combination product or kit:
 _____ Chart Note Page #: _____

UTAH DEPARTMENT OF HEALTH, PRIOR AUTHORIZATION REQUEST FORM

Dose, Age and/or Quantity Limits Exception Criteria for Approval:

Off label use criterion may apply. Medications with Clinical PA forms such as; Opioids, Buprenorphine Products, Antipsychotics in Children, etc. must be submitted on respective [Clinical PA forms](#)

- ☐ Member has failed to achieve adequate response within Medicaid's **Quantity/Dose Limit**.
Medication and dose: _____ Chart Note Page #: _____
Details of failure: _____
- ☐ Appropriate clinical rationale for prescribing medication outside Medicaid's **Age Limit**:
_____ Chart Note Page #: _____

Off Label or Compendia Use of FDA-Approved Drugs Criteria for Approval:

- ☐ Requests for any off-label indications must be supported by at least one (1) major multi-site study or three (3) smaller studies published in JAMA, NEJM, Lancet or other peer review specialty medical journals within the most recent five (5) years. Supporting documentation must be included. Compendia use must be recommended by generally-accepted compendia such as American Hospital Formulary Service Drug Information (AHFS), United States Pharmacopeia-Drug Information (USP-DI), and the DRUGDEX Information System.
- Diagnosis: _____
- Duration of treatment: _____

Re-authorization Criteria:

Updated letter with medical justification or updated chart notes demonstrating positive clinical response.

Authorization: Up to Six (6) months

Re-authorization: Up to one (1) year

PROVIDER CERTIFICATION

I hereby certify this treatment is indicated, necessary and meets the guidelines for use.

Prescriber's Signature

Date